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64 Enhancement of intranasal absorption of calcitonin by formulation with surfactants.

57 A pharmaceutical composition for the treatment of disorders of bone metabolism which comprises an aqueous or non-aqueous medium suitable for intranasal administration and containing a therapeutically effective amount of calcitonin and a surface active agent.

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1 ENHANCEMENT OF INTRANSAL ABSORPTION OF
 CALCITONIN BY FORMULATION WITH SURFACTANTS

5 The present invention relates to a novel method of
administering calcitonin to patients and to formulations
adapted for nasal administration.

10 Calcitonin is a polypeptide hormone isolated from
different organs in different species, including man and
salmon, or obtained via synthetic routes. Calcitonin is
15 recognized as being effective in diminishing hypercalcemia
and decreasing plasma phosphate concentrations in patients
with hyperparathyroidism, idiopathic hypercalcemia of
infancy, vitamin D intoxication, and osteolytic bone
metastases. While direct renal effects and actions on the
20 gastrointestinal tract are recognized, calcitonin is best
known for its effect on bone. Its use has proved to be
effective in diseases characterized by increased skeletal
resorption and abnormal bone formation, such as occurs for
example, in Paget's disease.

25 The method of administration of calcitonin is
predominantly by injection, although efforts were made in the
prior art to use other modes of administration, especially
for the treatment of localized conditions. While injectable
administration by physicians of calcitonin is proper for
30 short-term therapy, administration of calcitonin by injection
to patients in need of long-term calcitonin therapy has a
serious problem. Not only is it costly to patients to have
physicians do the administration of calcitonin for extended
periods of time but it is also painful and inconvenient. Nor

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1 can calcitonin be given orally to patients as it will be
destroyed by the digestive juices in the gastrointestinal
tract.

5 In view of the foregoing, it is apparent that a
serious need exists for a different route of delivery of
calcitonin to patients suffering from conditions that require
prolonged calcitonin therapy.

Nasal preparations are known in the prior art.
Generally, nasal preparations comprise an oil-in-water or
10 water-in-oil emulsion or an oily solvent base suitable for
use on the mucous membranes, such as mineral or vegetable
oils and fatty acid esters and one or more chemicals which
are soluble in the base. Such preparations usually contain
one or more active drugs intended to alleviate or mitigate a
15 condition in the body by their adsorption into the blood
stream through the mucous membrane of the nose.

While small molecules such as propranolol are
efficiently absorbed intranasally, large molecules such as
calcitonin show little if any absorption. The purpose of
20 this invention is to find agents capable of increasing the
bioavailability of calcitonin so that cost of therapy is
reasonable. The prior art has also recognized that the nasal
absorption of certain drugs may be facilitated by the use of
surfactants in such nasal preparation. For example, insulin
25 and polypeptides were found to have an improved absorption
rate when used in a solution containing a surfactant.

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1 It has now been found that hypercalcemia, Paget's
disease and other disorders of bone metabolism can be
advantageously treated by intranasal application of
calcitonin contain in a nasal preparation having an
5 absorption promoter and a buffer as essential ingredients.
Such preparations possess enhanced absorption across the
nasal mucosa when applied intranasally, but causes no
irritation or discomfort on extended use.

10 The present invention relates to a method for the
treatment of a mammal suffering from a disorder characterized
by high serum calcium which comprises intranasal application
of a nasal preparation containing a peptide having calcitonin
activity and an absorption promoting agent to effect control
of said disorders by transepithelial action.

15 According to the invention, calcitonin is
intranasally administered to a mammal via a novel dosage
form, such as a solution, ointment, or gel.

20 Calcitonin is a peptide hormone of 32 amino acids
with a disulfide bond at 1-7 in the amino terminus of the
molecule. These first seven amino acids with the disulfide
bond seem essential for activity and this sequence is
preserved from species to species. Calcitonin, as used
herein, means not only peptides having a structure
corresponding to one of the naturally occurring hormones, and
25 which may be naturally or synthetically produced, but also
related peptides having calcitonin activity.

30 The amount of calcitonin contained in the
preparation of the present invention may vary according to
various parameters, such as the nature of the preparation,

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1 the particular kind or activity of calcitonin employed and
the condition or ailment to be treated with the preparations.
In general, the concentrations are somewhat higher than those
found in compositions for the systemic administration of
5 calcitonin. It has been found that a concentration level of
1 to 150 micrograms per ml and preferably 2 to 30 micrograms
per ml achieve the desired result. The levels of
administration of calcitonin also vary somewhat from those
used systemically. In the case of human patients, for
10 example, amounts of from 0.7 to 70 micrograms, particularly
from 1 to 25 micrograms, are usually appropriate for single
dosages given and repeated as often as the physician finds it
necessary and such dosages correspond generally to about 0.01
to 1 micrograms, and particularly 0.03 to 0.35 micrograms,
15 per kilogram of body weight. (The above concentration and
dosage levels of calcitonin apply to calcitonin with a
potency of about 4000 International Units per mg and may be
adjusted pro rata for calcitonin of other potencies.)

The diluent base or vehicle used in accordance with
20 the present invention may be non-aqueous or aqueous. In the
former case the group of diluents is the physiologically
acceptable polar solvents. Preferred compounds of this type
are those with which it is possible to make a solution of
adequate concentration of dissolved calcitonin. Examples of
25 these compounds include dimethylsulphoxide, dimethyl
foramide, dimethylauramide, polyhydroxy alcohols, vegetable
and mineral oils. If desired, such non-aqueous media may be
mixed with water to form the diluent of the preparation.

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1 However, the degree of physiological acceptability of the
non-aqueous diluents is generally less than that of aqueous
media and the preferred diluent is therefore water without
the addition of organic solvents.

5 In the preparations of the present invention,
calcitonin is used in combination with an absorption
promoter. Such absorption promoters include the
physiologically acceptable surface active agents. The amount
of such an agent may be in the range from about 0.01 to about
10 10% w/v or higher and preferably about 0.05 to about 1.0%
w/v, the amount depending on the specific surfactant used.
The amount is generally kept as low as possible since above a
certain level no further enhancement of absorption can be
achieved and also too high of a surfactant level may cause
15 irritation of the nasal mucosa. Such surface active agents
include:

- a. Bile salts, such as sodium taurocholate, sodium cholate,
sodium deoxycholate and sodium glycholate;
- b. Cationics, such as the long chain amine condensates with
20 ethylene oxide and quaternary ammonium compounds, for example
cetyl trimethyl ammonium bromide and dodecyl dimethyl
ammonium bromide;
- c. Anionics, such as alkylbenzenesulfonates,
N-acyl-n-alkyltaurates, α -olefin sulfonates, sulfated
25 linear primary alcohols and sulfated polyoxyethylenated
straight-chain alcohols;
- d. Nonionics, such as polyoxyethylenated alkylphenols,
polyoxyethylenated straight chain alcohols, long chain
carboxylic acid esters including glycerol ester of natural
30 fatty acids, propylene glycol, sorbitol, and
polyoxyethylenated sorbitol esters;

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- 1 e. Amphoterics, such as imidazoline carboxylates,
sulfonates and the like; and
f. Phospholipids, such as phosphotidyl choline and the like.

5 The preparations of the present invention
preferably contain a phosphate or acetate buffer in the range
of 0.01 M to 0.5 M and preferably in the range of 0.05 M to
0.2 M. This concentration was found effective to provide
stability of the dissolved calcitonin in the diluent base or
vehicle.

10 The preparations of the present invention may also
contain other additives, such antioxidants, stabilizers,
tonicity adjusters, viscosity builders, preservatives, and
the like. The concentration of these additives may vary
according to the particular additive used and the desired
15 result sought. In general, the concentrations for these
additives will be in the range as follows:

	<u>Additives</u>	<u>% W/V</u>
	Antioxidants	0.01 - 0.2
	Stabilizers	0.01 - 2.0
20	Tonicity Adjuster	0.01 - 0.5
	Viscosity Builders	0.1 - 2.0
	Preservatives	0.001 - 2.0

25 While the use of the kind and concentration of
additives will be well within the ability of the skilled
artisan, the following will serve as illustration for two
additives generally used in pharmaceutical preparations
intended for similar purposes.

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1	<u>Preservatives</u>	<u>% W/V</u>
	Benzalkonium chloride	0.004 - 0.02
	Disodium Ethylene	
	Diamine Tetraacetate	0.01 - 0.2
5	Thimerosal	0.001 - 0.01
	Chlorobutanol	0.5 - 1.0
	Methyl and/or Propyl	
	Paraben	0.01 - 0.2
	Phenethyl Alcohol	0.25 - 0.75
10	Cyclohexedine	0.01 - 0.1

	<u>Viscosity Agents</u>	<u>% W/V</u>
	Methyl Cellulose	0.1 - 2.0
	Hydroxyethyl Cellulose	0.1 - 2.0
15	Hydroxypropyl Cellulose	0.1 - 2.0
	Polyvinylprrolidone	0.5 - 2.0

20 In preparing the formulations of the present invention, calcitonin is dissolved in the vehicle or diluent after which the additional ingredients are added in accordance with customary formulation procedures known in the pharmaceutical industry.

25 Examples of typical intranasal formulations are set forth below. However, it is to be understood that these examples are given by way of illustration only and are not to be construed as limiting the invention either in spirit or in scope as many modifications will be apparent to those skilled in the art.

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EXAMPLE 1% W/V

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Calcitonin	0.009
Sodium Taurocholate	0.5
Gelatin	1.0
Purified Water Q.S.	100

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EXAMPLE 2% W/V

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Calcitonin	0.009
Miranol C2M	1.0
Gelatin	1.0
Purified Water Q.S.	100

EXAMPLE 3% W/V

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Calcitonin	0.009
Miranol C2M	0.05
Sodium Acetate $.3H_2O$	1.36
Acetic Acid	0.6
Purified Water Q.S.	100

25

EXAMPLE 4% W/V

30

Calcitonin	0.009
Polysorbate 80	1.0
Sodium Acetate $.3H_2O$	1.36
Acetic Acid	0.6
Purified Water Q.S.	100

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EXAMPLE 5

% W/V

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Calcitonin	0.003
Brij 30	1.0
Sodium Acetate .3H ₂ O	1.36
Acetic Acid	0.6
Purified Water Q.S.	100

10

EXAMPLE 6

% W/V

15

Calcitonin	0.009
Myrj 59	1.0
Sodium Acetate	1.36
Acetic Acid	0.6
Purified Water Q.S.	100

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EXAMPLE 7

% W/V

Calcitonin	0.009
Miranol C2M	1.0
Sodium Phosphate	2.40
Citric Acid	0.34
Thimerasol	0.002
Purified Water Q.W.	100

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EXAMPLE 8

% W/V

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Calcitonin	0.009
Sodium Taurocholate	0.5
Sodium Acetate .3H ₂ O	1.36
Acetic Acid	0.6
Benzalkonium Chloride	0.01
DiSodium ethylenediame	
tetraacetate	0.1
Purified Water Q.S.	100

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EXAMPLE 9

% W/V

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Calcitonin	0.009
Sodium Taurocholate	0.5
Sodium Acetate .3H ₂ O	1.36
Acetic Acid	1.36
Chlorobutanol	0.1
Phenethyl Alcohol	0.2
Purified Water Q.S.	100

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EXAMPLE 10

% W/V

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Calcitonin	0.003
Miranol C2M	1.0
Sodium Phosphate	2.40
Citric Acid	0.34
Thimerasol	0.002
Purified Water Q.S.	100

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1 The gelatin used in the above formulations is a
standard hydrolipid animal gelatin prepared for
pharmaceutical use and routinely used as a diluent for
peptides.

5 According to the present invention, it has been
found that calcitonin can be administered intranasally from a
vehicle containing absorption promoters with results
considerably superior to those obtained with the
administration of calcitonin without absorption promoters.
10 The following studies were undertaken to examine the
bioavailability of calcitonin from the formulations of the
presetn invention, dependency of intranasal absorption of
calcitonin on the level of absorption promoters and stability
of calcitonin in the presence of absorption promoters.

15 PROTOCOL

Male rats weighing 150-250 g were weighed and
anesthetized with sodium pentobarbital, 50/mg/kg. by
intraperitoneal injection. Once anesthetized the
nasopalatine process was occluded with glue. The animals
20 were randomly placed into groups of 5-7 rats with the number
of groups being dependent upon the number of intranasal
formulations to be tested. Supplemental pentobarbital
anesthesia was administered as necessary throughout the
study.

25 Prior to administration of the test material, blood
was collected by cardiac puncture using a 25G 5/8" needle.
Fifty (50) microliters of the salmon calcitonin-containing
surfactant solution was then instilled into the nasal septum
using polyethylene tubing (PE 20, Peterson Technics, Monmouth
30 Junction, N.J.) connected to a 1 ml syringe; the tubing was
inserted about 1 cm into the nasal septum. One and three

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1 hours after nasal instillation, blood was again collected by
cardiac puncture.

Biochemical Analysis

5 Blood samples were allowed to clot at room
temperature and were then refrigerated for 30-60 minutes to
provide maximum clot retraction. The samples were
centrifuged at 4°C., 5000 rpm for 10 minutes (Beckman Model
J2-21 Centrifuge, Beckman Instruments, Palo Alto, CA). Serum
calcium was quantitated using a Calcette (Model 4008,
10 Precision Systems, Sudbury, MA).

Data Analysis

15 Serum calcium values at 0, 1 and 3 hours were
expressed as mean \pm standard deviation. In addition, the
absolute change and the percent change from the pretreatment
(0 time) value at 1 and 3 hours was also calculated.
Statistical analysis consisted of comparison of the serum
calcium values at 0 and 1 hour, 0 and 3 hours, and 1 and 3
hours using a t test.

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EXAMPLE 11

This example illustrates decrease in serum calcium in blood samples obtained in accordance with the above protocol when: a. calcitonin is administered alone;

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b. calcitonin is administered in formulations containing various absorption promoters; and c. no calcitonin is present in the formulations.

Table 1 shows the result obtained.

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TABLE I

Calcitonin U/kg body weight vehicle/surfactant	TIME AFTER DOSE			
	0 hour mg/dl	1 hour mg/dl	% decr.	3 hour mg/dl % decrease
2U 1% gel -	8.8	9.5	NONE	9.9 NONE
5U 1% gel -	8.5	7.9	7.1	9.5 NONE
10U 1% gel -	9.2	7.6	17.4	9.7 NONE
10U .1M Acetate -	8.8	6.3	28.4	9.0 NONE
- 1% gel 1% Miranol C2M(1)	8.9	8.8	NONE	9.2 NONE
- 1% gel 1% Taurocholate	9.1	9.5	NONE	9.8 NONE
3U 1% gel 1% Miranol C2M(1)	8.9	6.7	24.7	7.6 14.6
	8.8	6.5	26.1	8.5 3.4
3U 0.1M Acet. 1% Miranol C2M(1)	8.7	6.8	21.8	8.9 2.3
10U 1% gel 1% Miranol C2M(1)	8.7	6.8	21.8	8.9 2.3
	9.5	6.1	35.8	8.8 7.4
10U 0.1M Acet. 1% Miranol C2M(1)	9.1	7.5	17.6	6.6 27.5
	8.9	7.1	20.2	7.1 20.2
3U 1% gel 1% Taurocholate	9.3	6.1	34.4	6.5 30.1
10U 1% gel 1% Taurocholate	9.2	6.8	26.1	8.4 8.7
	9.1	7.0	23.1	6.4 29.6
10U 1% gel 1% Taurocholate	9.3	7.1	23.7	6.3 32.3
	8.6	6.1	29.1	5.9 31.4
10U 0.1M Acet. 1% Taurocholate	9.1	6.5	28.6	5.7 37.4
3U 1% gel 1% Tween 80 (Polysorbate 80)	8.3	6.2	25.3	8.7 NONE
	8.9	7.1	20.2	9.2 NONE

(1) Dicarboxylated fatty imidazoline or dicarboxylic coconut derivative,

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TABLE I (Cont'd.)

Calcitonin U/kg body weight vehicle/surfactant	0 hour mg/dl	TIME AFTER DOSE			3 hour mg/dl	% decrease
		1 hour mg/dl	% decr.			
10U 1% gel 1% Tween 80 (Polysorbate)	8.7	6.6	24.2	7.3	16.1	
3U 1% gel 0.5% Benzal- konium Chloride	8.9	6.7	24.7	8.4	5.6	
	8.7	5.9	32.2	6.3	27.6	
10U 1% gel 0.5% Benzal- konium Chloride	8.9	6.5	26.9	9.0	NONE	
	8.5	7.2	15.3	7.9	7.1	
3U 1% gel 1% Saponin (Sapogin Glycoside)	9.0	6.0	33.3	6.2	31.0	
	8.6	7.3	15.1	7.3	15.1	
10U .1M Acet. 1% NaL Saf	8.7	6.5	25.9	8.1	6.9	
10U .1M Acet. 1% Brifj 30	8.7	6.5	25.9	6.5	25.9	
(Polyoxyethylene (4) lauryl ether)						
10U .1M Acet. 1% Myrj 59	8.5	6.2	27.1	8.4	1.2	
(Polyoxyethylene (100) Stearate)						
10U .1M Acet. 1% Tween 80	8.7	7.5	13.8	7.1	18.4	
10U .1M Acet. 1% Aer OT	9.1	6.6	27.5	7.6	16.5	
(Sodium dioctyl sulfosuccinate)						

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EXAMPLE 12

This example illustrates that the enhancement of intranasal absorption depends on the level of absorption promoter present in the formulation.

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Table II shows the result obtained.

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TABLE II

10U Calcitonin/kilo in 0.1M Acetate with	0 hour mg/dl	1 hour mg/dl	3 hour mg/dl	%
1% Taurocholate	9.1	6.5	5.7	37.4
0.5%	9.0	6.1	7.5	16.6
0.25%	9.1	6.8	7.4	18.2
0.1%	8.9	6.5	8.3	6.7
0.05%	9.0	7.6	8.7	3.3

10U calcitonin/kilo
in 0.1M Acetate with

1% Miranol C2M (dicarboxylic coconut derivative, sodium salt)

0.5% "
0.25% "
0.1% "
0.5% "

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EXAMPLE 13

This example illustrates that calcitonin maintains its activity level in the formulations of the present invention on storage at room temperatures.

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Table III shows the results obtained.

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TABLE III

10U calcitonin in 1% gel with 1% Miranol C2M			
	0 hour mg/dl	1 hour mg/dl	%
initial	8.9	7.1	20.2
2 wks @ RT	9.1	7.2	20.9
4 wks @ RT	9.1	6.0	34.1
10U calcitonin in 1% gel with 1% Tween 80 (Polysorbate 80)			
	0 hour mg/dl	1 hour mg/dl	%
initial	8.9	6.7	24.7
2 wks @ RT	8.8	6.7	23.9
4 wks @ RT	8.8	6.3	28.4
	3 hour mg/dl		%
	7.1		20.2
	6.4		29.7
	6.9		24.2
	8.4		5.6
	8.3		6.0
	6.7		23.9

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1 What is claimed is:

1. A pharmaceutical composition for the treatment of disorders of bone metabolism which comprises an aqueous or non-aqueous medium suitable for intranasal administration and
5 containinng a therapeutically effective amount of calcitonin and a surface active agent.

2. The pharmaceutical composition of claim 1 further comprising a buffer.

3. The pharmaceutical composition of claim 2 wherein
10 said buffer is from 0.01 to 0.5M.

4. The pharmaceutical composition of any of claims 1-3 further comprising an antioxidant, stabilizer, tonicity adjuster, viscosity builder, or a preservative.

5. The pharmaceutical composition of any of claims 1-4
15 wherein said medium contains from about 5 to about 150 micrograms calcitonin per ml of said aqueous medium.

6. The pharmaceutical composition of any of claims 1-5 containing from about 0.01 to about 10% w/v of the surface active agent.

7. The pharmaceutical composition of claim 2 wherein
20 said buffer is a phosphate buffer.

8. The pharmaceutical composition of claim 2 wherein said buffer is an acetate buffer.

9. The pharmaceutical composition of any of claims 1-8
25 wherein said aqueous medium is a gel.

10. The pharmaceutical composition of any of claims 1-9 wherein said surface active agent is a dicarboxylated fatty imidazoline or sodium taurocholate, or a benzalkonium chloride.

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European Patent
Office

EUROPEAN SEARCH REPORT

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Application number

DOCUMENTS CONSIDERED TO BE RELEVANT			EP 83113070.3
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 7)
A	<u>US - A - 4 241 051</u> (CHRISTIE et al.) * Claims 3,4,5,7-10; column 1, line 45 - column 4, line 46 * --	1,6	A 61 K 37/02
A	<u>GB - A - 1 548 984</u> (CIBA-GEIGY AG) * Claims, especially claim 5; page 1, line 9 - page 3, line 66 * --	1,2,5	
A	<u>DE - A1 - 2 254 061</u> (HOECHST AG) * Claim 1,4; pages 1,2 * ----		
			TECHNICAL FIELDS SEARCHED (Int. Cl. 7)
			A 61 K 37/00
The present search report has been drawn up for all claims			
Place of search VIENNA		Date of completion of the search 30-03-1984	Examiner STÖCKLMAYER
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	